

Diversey Europe Operations BV-

Maarssenbroeksedijk 2 3542 DN Utrecht The Netherlands

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CERTIFICATE OF ANALYSIS

Date: 17_05.2017

Product Name	CLEARKLENS CLEANSINALD SS VH9S 100848254		
Product Code			
Batch Number	FMP 17187 42770		
Production Date	17,05,2017		
Expiration Date	EXP IT II I DOIS		

This is to certify that the above batch of product has been tested and conforms to the manufacturing Quality Control specification for the product.

Test	Test Method	Lin	nits	Results
Anna		Lower -	Upper	
Appearance	Visual	Clear Slig	htly Yellow	clear slightly
			uid	Yellow havid
pH (100% 20°C)	JDM001	9.0	11.4	11.0
Specific Gravity (20°C)	JDM004	0.990	1.010	0.996
	-(1)			0.110

On behalf of Diversey site Quality Manager	Name:	7. Stavon Olpa Linchne
	Position	Quality Control Inspector

inis accument being issued electro	nically does not bear a signature	
COA 100848254		
	Version: 02	Date of issuing: April 7th 2016



http://www.synergyhealthpic.com

Certificate of Irradiation

Date Issued: 08-May-2017

UK33S11867319-1-3

This is to certify that AST Daventry Synergy Health PLC has where appropriate delivered an irradiation process in accordance with:

EN ISO 11137-1:2015 Sterilisation of Health Care Products EN ISO 9001:2008 Quality Management System EN ISO 13485:2012 Quality System - Medical Devices

Flexible Medical Packaging Ltd Unit 8, White Cross Ind Estate Hightown Lancaster Lancashire LA1 4XS UNITED KINGDOM

All in accordance with current Technical Agreement

Order Information **Account Number:** 100432 Synergy Health Sales Part Reference: 1105380 **Customer Reference Number:** P028853 **Product Description:** C/SINALD SS 900ml Bottle DV4673 25-40kGy Validation Reference: 4673 **Quantity Received:** Customer Minimum Specification kGy: 25.0 **Customer Maximum Specification kGy:** 40.0 Customer Unit Lot/Batch Number: W042771, 1 PLT **Irradiation Data** Date and Time of Irradiation: 08-May-2017 03:13 Reference Dose Range kGy: 32.8 - 32.8 Calculated Minimum Dose kGy: 30.1 Calculated Maximum Dose kGy: 35.9

Items irradiated under WO42771 are to be used in finished product batch FMP17137 42770



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Ms D Henderson

Flexible Medical Packaging Limited

Unit 8 Hightown

White Cross Industrial Estate

Lancaster LA14XS

Date Received:

22 May 2017

Date Tested:

26 May 2017

Date Test Completed:

09 Jun 2017

Purchase Order:

28983

CERTIFICATE OF ANALYSIS

Laboratory Reference Number:

0035702/1

Test Required:

Sterility by Membrane Filtration Steritest

Date Received:

22/05/2017

Test Article:

ClearKlens Cleansinald SS

Sample Code:

FMP 17137 42770

Batch Ref:

8469

Qty Received:

20 x 900mL Bottles

Test	Method Item	Result
Sterility Test by Membrane Filtration (Steritest) Method	MM107/00	Pass
Growth in Tryptone Soya Broth at 20- 25°C after 14 days	MM107/01	No growth in one broth
Growth in Fluid Thioglycollate Medium at 30-35°C after 14 days	MM107/02	No growth in one broth
Volume Tested	MM107/05	20 x 50 mL
Tested in Accordance with Ph Eur, USP & JP	MQ005/07	EP 9.0 2.6.1, USP 40 <71> & JP XVII 4.06
Product Standard Data Sheet	FG047/psd	FM06

Comments

The testing procedure was performed in compliance with the current Pharmacopoeias however, the volume tested does not comply with the requirements of the current Pharmacopoeias.

Approval is provided by Electronic Signature. Their name and position is shown below.

Share Charles BSC CSIck MRSB

Date: 09 Jun 2017 18:03:41

Mrs S.L. Wood

Senior Laboratory Manager - Pharmaceutical Microbiology

Certificate of Analysis - OSMM Consignment: 0035702 Print Number: P0043410 Page 1 of 1

